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Novel peptides for use in immunotherapy of autoimmune diseases

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The invention relates to the use of novel peptides in a peptide induced tolerance therapy to prevent autoimmune disorders and in particular their use in treatment of chronic destruction of articular cartilage. The invention furthermore embraces pharmaceutical compositions comprising said peptides and a diagnostic method for the detection of autoreactive T cells in a test sample.

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NOVEL PEPTIDES FOR USE IN IMMUNOTHERAPY OF AUTOIMMUNE DISEASES

The invention relates to novel peptides, their use in treatment of chronic destruction of articular cartilag autoimmune diseases, pharmaceutical compositions comprising said peptide, and a diagnostic method f the detection of autoreactive T cells in a test sample.

The immune system is established on a principle of discrimination between foreign antigens (non-self antigens) and autoantigens (self antigens, derived from the individuals own body) achieved by a build-i tolerance against the autoantigens.

The immune system protects individuals against foreign antigens and responds to exposure to a foreign antigen by activating specific cells such as T- and B lymphocytes and producing soluble factors like

interleukins, antibodies and complement factors. The antigen to which the immune system responds is degraded by the antigen presenting cells(APCs) and a fragment of the antigen is expressed on the cell surface associated with a major histocompatibility complex (MHC) classII glycoprotein. The MHCglycoprotein-antigen-fragment complex is presented to a T cell which by virtue of its T cell recept recognizes the antigen fragment conjointly with the MHC classII protein to which it is bound. The T cel becomes activated, i. e. proliferates and/or produces interleukins, resulting in the expansion of the activ lymphocytes directed to the antigen under attack (Grey et al. , Sci. Am., 261 : 38-46, 1989).

Self antigens are also continuously processed and presented as antigen fragments by the MHC glycoproteins to T cells (Jardetsky etal., Nature211 : 326-329, 1991). Self recognition thus is intrinsic tc the immune system. Under normal circumstances the immune system is tolerant to self antigens and activation of the immune response by these self antigens is avoided.

When tolerance to self antigens is lost, the immune system may become activated against one or more self antigens, resulting in the activation of autoreactive T cells and the production of autoantibodies. This phenomenon is referred to as autoimmunity. As the immune response in general is destructive, i. e. meant to destroy the invasive foreign antigen, autoimmune responses can cause destruction of the body's own tissue.

The contribution of T cells to autoimmune diseases has been established in several studies. In mice, experimental autoimmune encephalomyelitis (EAE) is mediated by a highly restricted group of T cells, linked by their specificity for a single epitope of myelin basic protein (MBP) complexed to an MHC cla II molecule. In the Lewis rat, a species with high susceptibility to various autoimmune diseases, disease has been shown to be mediated by T cells. In humans autoimmune diseases are also thought to be associated with the development of auto-aggressive T cells.

A destructive autoimmune response has been implicated in various diseases such as rheumatoid arthritis (RA), in which the integrity of articular cartilage is destroyed by a chronic inflammatory process resulti from the presence of large numbers of activated lymphocytes and MHC class II expressing cells. The m presence of cartilage appears necessary for sustaining the local inflammatory response : it has been suggested that cartilage degradation is associated with the activity of cartilage-responsive autoreactive T cells inRA (Sigall et al. , Clin. Exp. Rheumat.~:59, 1988; Glant et al. , Biochem. Soc.

Trans. 18:796, 1990; Burmester etal., Rheumatoid arthritis Smolen, Kalden, Maini (Eds) Springer-Verla Berlin Heidelberg, 1992). Furthermore, removal of cartilage from RA patients by surgery was shown to reduce the inflammatory process(R. S. Laskin, J.

Bone Joint Surgery (Am) 72:529, 1990). The cartilage proteins are therefore considered to be target autoantigens which are competent of stimulating T cells. Activation of these autoreactive T cells leads t development of autoimmune disease. However, the identification of the autoantigenic components that play a role in the onset of rheumatoid arthritis has so far remained elusive.

The inflammatory response resulting in the destruction of the cartilage can be treated by several drugs, such as for example steroid drugs. However, these drugs are often immunosuppressive drugs that are nonspecific and have toxic side effects. The disadvantages of nonspecific immunosuppression makes this a highly unfavourable therapy.

The antigen-specific, nontoxic immunosuppression therapy provides a very attractive alternative for the nonspecific immunosuppression. This antigen-specific therapy involves the treatment of patients with the target autoantigen or with synthetic T

cell-reactive peptides derived from the autoantigen. These synthetic peptides correspond to T cell epitop of the autoantigen and can be used to induce specific T cell tolerance both to themselves and to the autoantigen. Although it seems paradoxical to desensitize the immune system with the very same antige responsible for activating the immune system, the controlled adminstration of the target (auto) antigen c be very effective in desensitization of the immune system. Desensitization or immunological tolerance c the immune system is based on the long-observed phenomenon that animals which have been fed or hav inhaled an antigen or epitope are less capable of developing a systemic immune response towards said antigen or epitope when said antigen or epitope is introduced via a systemic route.

The human cartilage glycoprotein-39 (HC gp-39) was previously identified as a target autoantigen in rheumatoid arthritis (RA) (Verheijden et al. , Arthitis Rheum.

4-Q : 1115-1125, 1997). The strategy followed for identification of relevant auto-epitopes within HC gp 39 was based on the assumption that the DR4 or DR1 molecules predispose to RA (Gao et al. , Arthitis Rheum.33 : 939-946, 1990 ; Nelson et al.,
Rheumatoid Arthritis, In Proceedings of the Eleventh International Histocompatibility
Workshop and Conference. Vol 1,Tsuji et al Ed, Oxford University Press, 1991) at two levels, firstly, by shaping the T cell repertoire and secondly, by determinant selection.

The shared epitope found among the RA-associated DR molecules might be involved in selection of similar sets of peptides for presentation to T cells (Gregerson et al. , Arthitis
Rheum.30:1205-1213, 1987). Putative binding sequences within the primary structure of HC gp-39 wer identified by use of a DR4(B1*0401) peptide binding motif (Verheijden et al. , Arthitis Rheum.40:1115 1125, 1997). HC gp-39, a protein of 362 amino acids, excluding the signal sequence (Hakala et al. , J. B Chem. 268 : 2580325810,1993), contains six regions accommodating this motif. Four peptides thus selected were synthesized and tested for binding the RA-associated DR1 and DR4(Bl*0401 and 0404) variants. All motif-based peptides, spanning residues 103-116, 259-275,263-275 and 326-338 of HC gp 39, were found to bind with high relative affinity to DRB 1 *0401 molecules. The recognition of these peptides by peripheral blood
T cells from RA patients and healthy donors was subsequently examined. All motif
based peptides were readily recognized in RA patients, thereby suggesting a high
frequency of HC gp-39-specific T cells in RA. The response to 263-275 was most prominent ; 8 out of 1 RA patients responded to this peptide (Verheijden et al. , Arthitis
Rheum.40:1115-1125, 1997). Thus, HC gp-39 is a target for immune recognition in the joint.

The significance of this protein for arthritic disease was further demonstrated by its arthritogenicity in Balb/c mice. A single injection in the chest region withHg amounts of protein mixed in IFA, induced a chronic joint inflammation reminiscent of RA (Verheijden et al. , Arthitis Rheum.40:1115-1125, 1997).

Recently, a novel human chondrocyte protein, YKL-39, was isolated and described (Hu et al.,J. Biol. Chem. 271 : 19415-19420,1996). The protein shares significant sequence identity with HC gp-39 (YKL 40). Another homologue of HC gp-39 is secreted by human macrophages and is termed chitotriosidase (Boot et al.,J. Biol. Chem.

270 : 26252-26256,1995). The sequences corresponding to the HC gp-39 (263-275) peptide RSFTLASSETGVG (SEQ IDN0 : 3) are identified as HSFTLASAETTVG (SEQ IDN0 : 2) within the YKL-39 protein (266-278) and as RSFTLASSSDTRVG (SEQ IDN0 : 4) within macrophage chitotriosidase (269-282) respectively (Table 1).

The chitotriosidase peptide Chi (269-282) contains theDRB1*0401 peptide binding motif which was previously used for selection of T-cell epitopes within proteins. In contrast, the YKL-39 (266-278) pepti

does not contain this 0401 motif.

It will be clear that tolerization of HC gp-39(263-275) -reactive T-cells may be of benefit to RA patients Likewise, mimicry epitopes of HC gp-39 (263-275) may have a similar function and may be used to induce tolerance. Preferably such mimicry epitopes will have at least the same tolerizing capacity.

To effectively use tolerance induction therapy to treat T cell mediated cartilage destruction, there is a gr need to identify T cell-reactive peptides which can desensitize patients against the autoantigen that is activating the T cells responsible for the inflammatory process.

Although the YKL-39 peptide does not contain the 0401 motif, it was surprisingly found that the YKL-39 (266-278) epitope is a mimicry epitope of HC gp-39 (263-275).

This epitope therefore is useful for tolerization of autoreactive T-cells with reactivity to HC gp-39 (263-275), YKL-39 (266-278) or their mimicry epitopes in rheumatoid arthritis patients.

It is an object of the invention to provide peptides which are able to induce systemic immunological tolerance, more in particular specific T cell tolerance, preferably to the responsible cartilage antigen in patients suffering from T cell-mediated cartilage destruction. The peptides of the present invention are characterized in that they comprise one or more of the amino acid sequences FTLASAETT (SEQ IDNO 1). More specifically, a peptide according to the invention comprises HSFTLASAETTVG (SEQ IDNO : 2).

Also within the scope of the invention are multimers of the peptides according to the invention such as f example a dimer or trimer of the peptides according to the invention. A multimer according to the invention can either be a homomer, consisting of a multitude of the same peptide, or a heteromer consisting of different peptides.

The characteristic amino acid sequences of the peptides according to the invention can be flanked by random amino acid sequences. Preferred are flanking sequences, that have a stabilizing effect on the peptides, thus increasing their biological availability.

Human Cartilage glycoprotein 39 is a target autoantigen in RA patients which activates specific T cells, thus causing or mediating the inflammatory process. HC gp-39 derived peptides were predominantly recognized by autoreactive T cells from RA patients but rarely by T cells from healthy donors, thus indicating that HC gp-39 is an autoantigen in RA. The arthritogenic nature of HC gp-39 was further substantiated in the
Balb/c mouse. A single, subcutaneous injection of said protein in Balb/c mice was able to initiate arthrit signs in the animals. The course of the HC gp-39- induced disease was characterized by relapses occuri periodically in fore paws and/or hind paws and gradually developed from a mild arthritis into a more severe form. Also, a symmetrical distribution of afflicted joints was observed which is, together with th observation of recurrent relapses, reminiscent of disease progression in arthritis, especially RA.

It was surprisingly found that the YKL-39 266-278 peptide was effective as a tolerogen. It will be clear those skilled in the art that the peptides may be extended at either side of the peptide or at both sides anc still exert the same immunological function. The extended part may be an amino acid sequence similar t the natural sequence of the proteinYKL-39.

The peptides according to the invention can be prepared by well known organic chemical methods for peptide synthesis such as, for example, solid-phase peptide

synthesis described for instance in J. Amer. Chem. Soc.$5 : 2149 (1963) and Int. J.

Peptide Protein Res.35 : 161-214 (1990). The peptides according to the invention can also be prepared b recombinant DNA techniques. A nucleic acid sequence coding for a peptide according to the invention (multimer of said peptides is inserted into an expression vector. Suitable expression vectors comprise the necessary control regions for replication and expression. The expression vector can be brought to expression in a host cell. Suitable host cells are, for instance, bacteria, yeast cells and mammalian cells. Such techniques are well known in the art, see for instance Sambrooke etal, MolecularCloning : a Laboratory
Manual, Cold Spring Harbor laboratory Press, Cold Spring Harbor, 1989.

The peptides may be stabilised by C- and/or N- terminal modifications, which will decrease exopeptidas catalysed hydrolysis. The modifications mayinclude : C-terminal acylation,(e. g. acetylation = Ac-peptic N-terminal amide introduction,(e. g. peptidNH2) combinations of acylation and amide introduction(e.g. Ac-peptide-NH2) and introduction of D-amino acids instead of L-amino acids (Powell et al. , J. Pharm. Sci. ,-U : 731-735, 1992).

Other modifications are focussed on the prevention of hydrolysis by endopeptidases. Examples of these modificationsare : introduction of D-amino acids instead of L-amino acids, modified amino acids, cyclisation within the peptide, introduction of modified peptide bonds,e. g. reduced peptide bonds [CH2NH] ande. g. peptoids (N-alkylated glycine derivatives) (Adang et al, Recl. Trav. Chim. Pays-Bas,113 : 63-78, 1994 and Simon et al, Proc. Natl. Acad. Sci. USA, 89:9367-9371, 1992).

The peptides according to the invention are T-cell epitopes, which are recognized by and are able to stimulate autoreactive T-cells. These autoreactive T cells may be founde. g. in the blood of patients suffering from autoimmune diseases.

Thus, according to the invention the peptides, said peptides resembling the MHC
Class II restricted T-cell epitopes present on the target autoantigen comprising the peptide of SEQ ID N' 1 or SEQ ID N0 : 2, are very suitable for use in a therapy to induce specific T-cell tolerance to said autoantigen in mammals, more specifically humans, suffering from T-cell mediated cartilage destructio such as for example arthritis, more specifically rheumatoid arthritis. Optionally such a treatment can be combined with the the administration of other medicaments such as DMARDs (Disease
Modifying Anti-Rheumatic Drugse. g. sulfasalazine, anti-malarials (chloroquine, hydroxychloroquine) injectable or oral gold, methotrexate, D-penicillamine, azathioprine, cyclosporine, mycophenolate), NSAIDs (non steroidal anti inflammatory drugs), corticosteroids or other drugs known to influence the course of the disease in autoimmune patients.

The peptides according to the invention can also be used to modulate lymphocytes that are reactive to antigens other than said autoantigen but are present in the same tissue as the autoantigen i. e. proteins or parts thereof comprising the peptide according to SEQ
IDN : l or SEQ IDN0 : 2. By the induction of antigen-specific T-cell tolerance, autoimmune disorders c be treated by bystander suppression. More in general, the cells to be modulated are hematopoietic cells. general, in order to function as a tolerogen the peptide must fulfill at least two conditions i. e. it must possess an immune modulating capacity and it must be expressed locally usually as part of a larger prot

Thus, the present invention provides a method to treat patients suffering from inflammatory autoimmun diseases, by administration of a pharmaceutical preparation comprising the peptide according to the invention. Such patients may suffer from diseases like Graves'diseases, juvenile arthritis, primary glomerulonephritis, osteoarthritis, Sjogren's syndrome, myasthenia gravis, rheumatoid arthritis, Addisor disease, primary biliary sclerosis, uveitis, systemic lupus erythematosis, inflammatory bowel disease,

multiple sclerosis or diabetes. The peptides according to the present invention therefore can be used in tl preparation of a pharmaceutical to induce tolerance in patients suffering from these diseases.

Treatment of autoimmune disorders with the peptides according to the invention makes use of the fact tl bystander suppression is induced to unrelated but co-localized antigens. The regulatory cells secrete in a antigen specific fashion pleiotropic proteins such as cytokines which may downmodulate the immune response.

According to the invention, patients suffering from T-cell mediated destruction of the articular cartilage can be treated with a therapeutical composition comprising one or more peptides according to the invention and a pharmaceutical acceptable carrier.

Administration of the pharmaceutical composition according to the invention will induce systemic immunological tolerance, in particular tolerance of the specific autoreactive T cells of these patients, to the autoantigenic proteins in the articular cartilage under attack and other self antigens which display the identified MHC Class II binding T cell epitopes characterized or mimicked by the amino acid sequences of one or more of the peptides according to the invention. The induced tolerance thus will lead to a reduction of the local inflammatory response in the articular cartilage under attack.

Very suitable peptides to be used in a pharmaceutical composition according to the invention are the peptides comprising the YKL-39 (268-276) or the YKL-39 (266-278) peptide flanked sequences up to a total length of 55 amino acids. More preferably the peptides have a length of 25 amin acids. Even more preferably the amino acid sequence of the peptides is FTLASAETT or HSFTLASAETTVG.

The peptides according to the invention have the advantage that they have a specific effect on the autoreactive T cells thus leaving the other components of the immune system intact as compared to the nonspecific suppressive effect of immunosuppressive drugs. Treatment with the peptides according to tl invention will be safe and no toxic side effects will occur.

Systemic immunological tolerance can be attained by administering high or low doses of peptides according to the invention. The amount of peptide will depend on the route of administration, the time o administration, the age of the patient as well as general health conditions and diet.

In general, a dosage of0. 01 to 10000ag of peptide per kg body weight, preferably0. 05 to 500ag, more preferably0. 1 to 100ug of peptide can be used.

Pharmaceutical acceptablecarriers are well known to those skilled in the art and include, for example, sterile saline, lactose, sucrose, calcium phosphate, gelatin, dextrin, agar, pectin, peanut oil, olive oil, sesame oil and water. Other carriers may be, for example MHC class II molecules, if desired embedded liposomes.

In addition the pharmaceutical composition according to the invention may comprise one or more adjuvants. Suitable adjuvants include, amongst others, aluminium hydroxide, aluminium phosphate, amphigen, tocophenols, monophosphenyl lipid A, muramyl dipeptide and saponins such as Quill A. Preferably, the adjuvants to be used in the tolerance therapy according to the invention are mucosal adjuvants such as the cholera toxine B-subunit or carbomers, which bind to the mucosal epithelium. The amount of adjuvant depends on the nature of the adjuvant itself.

Furthermore the pharmaceutical composition according to the invention may

comprise one or more stabilizers such as, for example, carbohydrates including sorbitol, mannitol, starch, sucrosedextrin and glucose, proteins such as albumin or casein, and buffers like alkalir phosphates.

Suitable administration routes aree. g. intramuscular injections, subcutaneous injections, intravenous injections or intraperitoneal injections, oral administration and nasal administration such as sprays.

It is another object of the invention to provide a method for detecting autoreactive T cells involved in th destruction of articular cartilage and test kits to be used in said method. Thus, the peptides according to invention are also very suitable for use in a diagnostic method to detect the presence of activated autoreactive T cells involved in the chronic inflammation and destruction of the articular cartilage.

The diagnostic method according to the invention comprises the following steps :
a) isolation of the peripheral blood mononuclear cells (PBMC) from a blood sample of an individual,
b) culture said PBMC under suitable conditions,
c) incubation of said PBMC culture in the presence of one or more peptides according to the invention, a
d) detection of a responseof T cells, for example a proliferative response, indicating the presence of activated autoreactive T cells in the individual.

The detection of a proliferative responseof T cells can be detected by, for example, the incorporation of thymidine.

Also within the scope of the invention are test kits which comprise one or more peptides according to th invention. These test kits are suitable for use in a diagnostic method according to the invention.

The following examples are illustrative for the invention and should in no way be interpreted as limiting the scope of the invention.

Legends to the figures
Figure1
Figure la, b, c. Cross reactivity of three, different,HC gp-39-specific hybridomas(8B12, 14G11, 20H5) with YKL-39 (266-278)
(CVR0271B = HC gp-39 (263-275), KV0432B = YKL-39 (266-278), CC0332B = Chi (269-282), KV0431A = YKL-39 (262-274).HCDA. 8B12. 1D8, 14G11. 1H7 and20H5.4F6.2F6 are HLA-DRB1 *0401 -restricted hybridomas specific for HC gp-39 (263275). Activation of T-cell hybridomas is expressed as IL-2 production.

Figure 2 In vivo tolerization with HC gp-39 (263-275) or YKL-39 (266-278)
Balb/c mice were tolerized by intranasal application of 50,10 or 2 microgram of HC gp-39 (263-275) or YKL-39 (266-278) followed by immunization with HC gp-39 (263-275). Mice t were pretreated with saline or that were left untreated were included as controls.

Examples
Example 1 Alignment of sequences
The human chondrocyte protein, YKL-39 shares significant sequence identity with HC gp-39 (YKL-40). Another homologue of HC gp-39 is secreted by human macrophages and is terme chitotriosidase (Boot et al. , 1995). The sequences corresponding to RSFTLASSETGVG (HC gp-39 (26 275), SEQ IDN0 : 3) were identified as HSFTLASAETTVG within the YKL-39 protein (266-278) and RSFTLASSSDTRVG (SEQ IDN0 : 4) within macrophage chitotriosidase (269-282) respectively (Table 1). Chi (269-282) contains theHLA-DRB1*0401 peptide binding motif which was previously used for selection of T-cell epitopes within proteins. In contrast, the YKL-39 (266-278) peptide does not contain

this motif. All peptides were synthesized.
Table 1. Alignment of the HC gp-39 (263-275) sequence with the corresponding region in YKL-39 and macrophage ChitotriosidaseHCgp-39 263-275 R S F T L A S S - E T G V G
YKL-39 266-278 H S F T L A S A - E T T V G
Chi (269-282)R S F T L A S S S D T R V G
Example 2 Binding of peptides toHLA-DRB1 *0401
The peptides from example 1 were tested for binding theDRB1*0401-encoded molecules. HLA-DR4 (DRB1*0401) molecules were purified from the homozygous
EBV-transformed human B lymphoblastoid cell linesHulyl38IC2 and the competition peptide HLA-DR binding assay was performed basically as described by Verheijden et al. , 1997. The affinity of a given peptide for binding DRB1*0401-encoded molecules was related to competition with a marker peptide. This relative binding affinity was defined as the peptide concentration at which the signal was reduced t 50% (IC50). The
HA-F peptide is a positive control (Hemagglutinin 307-319; PKFVKQNTLKLAT; at position 309 Y is substituted byF ; SEQ IDN:5). The peptide is known to have a high affinity for DRB 1 *0401 molecules

As expected, theChi (269-282) peptide was found to bind with high affintiy toDRB1*0401 (see table 2). The YKL-39 (266-278) peptide, which does not accommodate the effectiveDRB1*0401 peptide binding motif, bound with very high affinity to DR4(B1*0401).

Table 2 Peptide binding to HLA-DRBI *0401-encoded molecules
EMI11.1

<tb> <SEP> IC50 <SEP> values
<tb> peptide <SEP> batch <SEP> Exp. <SEP> A <SEP> Exp.B <SEP> Exp. <SEP> C
<tb> <SEP> YKL39(262-274) <SEP> KV0431A <SEP> 0.006 <SEP> 0.005 <SEP> ND <SEP>
<tb> <SEP> YKL39(266-278)KV432B0.0350.0320.12
<tb> <SEP> HCgp39(263-275) <SEP> CVR271B <SEP> ND <SEP> 0.008 <SEP> 0.038 <SEP>
<tb> <SEP> Chi(269-282) <SEP> CC0332B <SEP> 0.053 <SEP> 0.11 <SEP> 0.16 <SEP>
<tb> HA-FAE0690A0. <SEP> 200. <SEP> 140. <SEP> 20
<tb>
ND = not determined
Example 3 Stimulation T-cell hybridomas
Hybridomas specific for HC gp-39 (263-275) were tested for recognition of the corresponding sequences.

To test the cross reactivity of the 3 different, HC gp-39-specific hybridoma cell lines with theYKL-39 or the chitotriosidase peptide, 5 x104 hybridoma cells and 2 x105 irradiated (120(RAD), EBV-transformed B cells carrying theDRB1*0401 specificity were incubated in150 pI volumes : wells of a round-bottomed microtiter plate. Peptide antigen (HC gp-39 (263-275), YKL-39 (266-278), chitotriosidase (269-282) or a control peptide) was added in 50 pI volumes to duplicate wells. Forty-eigl hr later 100pLI of the culture supernatant was assayed for IL-2 production using a sandwich ELISA withPharmingen antibodies specific for mouse IL-2.

It was found that the synthetic peptide YKL-39 (266-278) generated a response similar to HC gp-39 (26 275) whereas the Chi (269-282) did not generate a response.

The data suggest that the three different TCRs utilized by three different hybridomas do not discriminat between HC gp-39 (263-275) or YKL-39 (268-278) when presented byDRB1*0401-encoded molecules (Figure la, b, c) but do discriminate between HCgp-39 (263-275) and Chi (269-282). The data indicate t

YKL-39 (266-278) is a mimicry epitope ofHC gp-39 (263-275). (Figla, b, c)
Example 4 Recognition of YKL-39 (266-278) by PBMC
Peripheral blood mononuclear cells (PBMC) were isolated from heparinized peripheral blood by standa centrifugation on Ficoll-Paque(Pharmacia, Uppsala,
Sweden). Cells were suspended in wells of a 24 well plate in a concentration of 5 x105 cells per ml. Cel were incubated in medium alone or in the presence of 10 or50 ,ug/ml peptide antigen (YKL-39 (266-278)). Cultures were incubated for 6 days at37 C in a humidified atmosphere of5% C02. Cells were the suspended and 100 or 150fil volumes of medium was distributed in 4-fold in wells of a 96-well roundbottomed plate.

Cells were then pulsed with
PBMC coincides with recognition of HC gp-39 (263-275) and HC gp-39 and
furthermore that recognition of YKL-39 (266-278) is generally more pronounced than
recognition of HC gp-39 (263-275).

Table 3a. Recognition of the YKL-39 (266-278) epitope by PBMC from RA patients.

Donor typingSI SI
10 llg/ml 50 ,ug/ml
242-0.2 NR 0404/15 3 < 2
337-0. 2 R 0401/02 19 58
338-0. 1 NR 03/14 < 2 < 2
454-0 R0401/9 9
456-0 R ND 15 4
457-0 NR ND < 2 < 2
458-0 R ND 4 27
459-0 R ND < 2 25
460-0 NR ND 3 < 2
SI = antigen-specificcounts/background counts. SIs 5 are regarded positive R = responder, NR = non-responder
Table 3b. Recognition of YKL-39 (266-278) coincides with recognition of HC gp
39 (263-275) and HC gp-39 protein.

Donor R/N YKL-39(266-278) HCgp-39 (263-275) HC gp-39 SI SI SI SI SI SI
Hg/ml 10 50 10 50 10 50
169 R 27 32 10 27 24 44
455 R 20 35 1 15 45 95
447 NR 12 1 1 1 1
327 R 6 5 3 5 12 19
SI = antigen-specific counts/background counts. SI > 5 are regarded positive. R = responder, NR = non-responder. NT = not tested. Donor 447 responds to Tetanus toxoid and Candida Albicans.

Example 5 Tolerance induction
A HC gp-39(263-275) -specific DTH assay suitable to monitor tolerance induction with peptide antigen was developed. Immunisation of Balb/c mice with HC gp-39 (263-275) in incomplete Freunds adjuvant (IFA) was found to be effective in the induction of a DTH response following challenge with the HC gp (263-275) peptide.

This peptide-based DTH system was used to detect modulation of the DTH response by nasal applicatio of HC gp-39 (263-275) peptide. It was found that nasal application of
HC gp-39 (263-275), in a dose-dependent manner, downmodulated the HC gp-39 (26275) -induced DTI

response. Nasal application of YKL-39 (266-278), however, resulted in a more enhanceddownmodulati of the DTH response, indicating that YKL-39 (266-278) can efficiently tolerize a peptide-specific respo: induced with HC gp-39 (263-275) (Table 4, Figure2a, b, c).

Table 4. Experimental set-up tolerization experiment
Pretreatment sensibilisation challenge tolerancenone HC gp-39 (263-275) HC gp-39 (263-275) no salin HC gp-39 (263-275) HC gp-39 (263-275) no
HC gp-39 (263-275) HC gp-39 (263-275) HC gp-39 (263-275) yes
YKL-39 (266-278) HC gp-39 (263-275) HC gp-39 (263-275) yes

Claims of corresponding document: **WO0005254**

Claims
1. Peptide having an amino acid sequence of 9-55 amino acid residues comprising the amino acid sequence FTLASAETT(SEQID NO:1).

2. Peptide according to claim1 comprising the amino acid sequence HSFTLASAETTVG (SEQ ID N0 :

3. Peptide according to claim1 or 2 having an amino acid sequence of up to 25 amino acid residues.

4. Peptide according to claim1 or 2 having the amino acid sequence FTLASAETT (SEQID NO:1) orHSFTLASAETTVG (SEQ ID N0 : 2).

5. Peptides according to any of the claims 1-4 for use as a therapeutical substance.

6. Pharmaceutical composition comprising one or more of the peptides according to claims1-4, and a pharmaceutical acceptable carrier.

7. Use of one or more of the peptides according to claims 1-4 for the manufacture of a pharmaceutical preparation for the induction of specific T-cell tolerance to an autoantigen in patients suffering from autoimmune disorders, more specifically arthritis.

8. Diagnostic composition comprising one or more of the peptides according to any of the claims 1-4 and a detection agent.